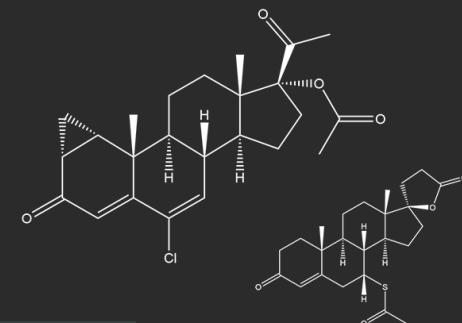


/HRTGen/

Male to Female Guide

Quickguide v2.51 (2022)



HRT OVERVIEW

Hormone Replacement Therapy (HRT) will change your life. The most common way to start is to take both an anti-androgen (AA) and an estrogen (E). The function of the AA is to tell cells to stop acting male, and the function of the estrogen is to tell cells to act female.

The AA is responsible for blocking testosterone (T) from acting on the androgen receptors (ARs) of cells. Some AAs will also damage testosterone production over time, and most but not all will result in lowering testosterone levels. The estrogen will activate estrogen receptors (ERs) in cells, which tells them to become female. When the brain detects higher levels of estrogen it will also start telling the body to produce less testosterone.

The combination of an estrogen and an anti-androgen can have tremendous feminizing effects. The most noticeable of these are skin softening, breast development, reduced body hair, stopping/reversing male pattern baldness, muscle atrophy, and redistribution of body fat into a female shape. These changes can make someone look exceedingly more feminine in the face and body, and in the worst cases only younger looking.

The results of HRT are also heavily dependent on the individual's health, genetics, and age. Different people can have vastly different results from the same treatment. Thus the goal of HRT is to optimize your dose and regimen to fit with your body's individual biology. Knowing the symptoms to watch for and taking blood tests regularly are crucial to ensuring the safest and most effective medical transition possible.

This guide is being provided in the interest of harm reduction, suicide prevention, and promoting responsible HRT. It is not intended to be a substitute for professional medical advice. When taking HRT it is advisable to be under the care of a physician you respect and trust, so that your health can be monitored and so you can get prompt treatment in case of any complications.

1. ANTI-ANDROGEN (AA)

a) CYPROTERONE ACETATE (Cypro, CPA)

Dosage: 5-12.5mg/day. Studies have shown 10mg to be effective as an AA (50-70% lowered T @10mg/day).

Action: Strong AA. Nukes T production and weakly blocks T.

Due diligence: Doses above 12.5mg are not recommended. Higher doses for prolonged periods are linked to venous thromboembolism and benign brain tumors. May raise prolactin levels. Monitor yourself for pseudo-lactation and monitor prolactin levels via blood tests. Higher doses of 100mg/day are associated with liver toxicity. Cypro is linked to vitamin B12 deficiency. To prevent B12 deficiency you should eat animal products, drink lots of milk, and/or take a B12 supplement to ensure healthy B12 levels.

Info: Effective in reducing general gonad function. Affordable AA.



b) BICALUTAMIDE (Bica)

Dosage: 25-100mg/day. 10-25mg/day will block female T levels which is achieved via estradiol at a sufficient dose.

If not taking estradiol or if you have male levels you will need 50mg/day or more. 100-150mg/day at minimum appears to be needed to fully or nearly-block 600ng/dL T.

Action: Androgen receptor antagonist (blocks androgens). Does not lower T but it does block its effects with sufficient dosages.

Due diligence: Avoid if you have a history of liver issues. Low incidence of liver level changes, and lower rates of liver toxicity. Issues present themselves in first 3-4 mo.

Info: Less effective at reducing general gonad function. More expensive AA. Buildup time for bica to 50% of steady levels is reached after 1 week, about 80-90% steady state levels after 3-4 weeks, and 100% after 6-12 weeks of continuous daily administration.



c) SPIRONOLACTONE (Spiron)

Dosage: 50-200mg/day. At doses of 50-200mg/day spiro is more effective at blocking female T levels levels and less suited at blocking male T levels.

Action: A potassium sparing diuretic that is a weak AA. Weakly blocks production of androgens, and blocks androgens.

Due diligence: Avoid potassium supplements. Avoid if you have kidney issues. Higher doses of spiro is associated with elevated cortisol levels (visceral adiposity), brain fog, and depression.

Info: Makes you pee a lot. Affordable AA. Off-target action with high antimineralocorticoid activity and mixed estrogenic and antiestrogenic or SERM-like activity (which has been implicated in having a negative impact on breast development).



d) GnRH AGONISTS (GnRH, GnRHs)

Types: Buserelin, Lupron (leuprolide), Goserelin, Triptorelin.

Action: Nukes LH/FSH, thus gonadal androgen production, and causes pituitary to be desensitized to GnRH (GnRH causes release of LH/FSH) with consistent use.

Info: Great alternatives if they can be obtained affordably. I.e. they are prohibitively expensive for most, and may be available through your GP/endo via insurance/public healthcare.



e) ESTRADIOL MONOTHERAPY (E MONO, E MONOTHERAPY)

Dosage: Estradiol (e2) levels of 200-500pg/ml suppress T levels by ~90%, and e2 levels between 200-500pg/ml suppress T levels by ~90-95%. This may vary due to capacity for gonads to produce androgens (e.g. T), and therefore your "monotherapy levels" may differ here (e.g. 90% suppression of 400ng/dL is "sufficient" as compared to 90% suppression of 700ng/dL).

Action: Lowers LH/SH via the brain registering it has sex hormones, and via lowering LH/FSH it tells the gonads to stop producing T.

Info: Easily and reliably attainable via injections. Can be attained via patches, gel, or sublingual.



2. ESTROGEN (E)

a) INJECTIONS

Types: Estradiol valerate 0.5mg, 1mg, 2mg, 5mg, 10mg, 15mg, 20mg. Estradiol cypionate 5mg. Estradiol enanthate 0.5mg, 1mg. Polyestradiol phosphate 0.5-2.5mg.

Dosing: Patches or gels. Patches come in reservoir or matrix forms. Changing patches: reservoir every 3.5 days, matrix every 7 days.

Dosing: Patches start at either 50-100ug, 150-200ug or more is the final dose per your goal levels & labs. 100ug patch is approximately equal to 1000pg/ml for levels. Transdermal gel starts at an equivalent dosage. Gel with 0.05% concentration has 1.5mg estradiol in 2.5g gel. Estradiol levels achieved with 1.5mg of estradiol gel are similar to those with a 500ug patch.

Info: Patches are known to leave visible residue after removal. Reservoir patches are known to cause skin reactions (~14.2% occurrence rate). Gel and transdermal application areas do change absorption, and smaller application areas for gel gives higher levels. Application area effectiveness: scrotum > buttocks > stomach > thigh = arm > hand = foot.

b) TRANSDERMAL

Types: Patches or gels. Patches come in reservoir or matrix forms. Changing patches: reservoir every 3.5 days, matrix every 7 days.

Dosing: Patches start at either 50-100ug, 150-200ug or more is the final dose per your goal levels & labs. 100ug patch is approximately equal to 1000pg/ml for levels. Transdermal gel starts at an equivalent dosage. Gel with 0.05% concentration has 1.5mg estradiol in 2.5g gel. Estradiol levels achieved with 1.5mg of estradiol gel are similar to those with a 500ug patch.

Info: Patches are known to leave visible residue after removal. Reservoir patches are known to cause skin reactions (~14.2% occurrence rate). Gel and transdermal application areas do change absorption, and smaller application areas for gel gives higher levels. Application area effectiveness: scrotum > buttocks > stomach > thigh = arm > hand = foot.

c) SUBLINGUAL (sub) & BUCCAL

Types: Estradiol hemihydrate (17-beta estradiol) or estradiol valerate (EV) pills can be used.

Dosing: Start at 2mg/day and later increase to 4-8mg/day. Split doses throughout the day as to maintain steadier levels. 2mg dosing example: 0.5mg/4x day (e.g. 16hr day, every 4 hrs take 0.5mg). It is recommended to split your dose >=3/day for more stable levels.

Info: Sublingual administration method is to dissolve under your tongue, and buccal is to dissolve between your cheek and gums. Do not eat or drink while they're dissolving. Wait 10-15min before eating or drinking after the pills have dissolved. Splitting or crushing the pills may help them dissolve faster. This is especially applicable to pills such as progynova (EV) due to the sugar coating.

d) ORAL

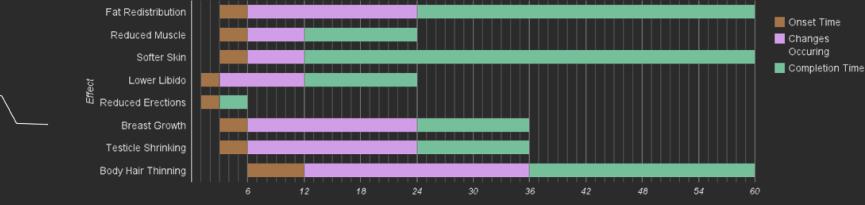
Types: Estradiol hemihydrate (17-beta estradiol) or estradiol valerate (EV) pills can be used. 2mg EV is roughly equivalent to 1.5mg estradiol hemihydrate.

Dosing: Start at 2mg/day and later increase to 4-8mg/day. Split doses throughout the day as to maintain steadier levels. 2mg dosing example: 0.5mg/4x day (e.g. 16hr day, every 4 hrs take 0.5mg). It is recommended to split your dose >=3/day for more stable levels.

Info: Oral administration method is to swallow the tablets. Significantly raises SHBG levels. 5-10mg/daily boron is recommended to counteract this. Due to lower e2 levels the main consideration when starting on oral e is T suppression. Cypro is more reliable here, and for bica it's recommended to get a pre-hrt blood test (FT) as to determine the proper dosage as to block T.

MTT HRT Effect Timeline

Note: highly variable. Depends on how quickly androgens are suppressed, and then E being the dominant sex hormone. I.e. can be much faster or slower.



3. QUICK DOSING GUIDE

MONOTHERAPY DOSING (INJECTIONS)

Use "injectable estradiol simulator" to refine dose (search it). Example 250pg/ml trough estradiol monotherapy dosages:

ESTER

ESTER	DOSAGE
EV	5mg/5days or 9mg/week
EC	5mg/week or 15mg/2 weeks
EEn	4mg/week or 14mg/2 weeks

ANTI-ANDROGEN DOSING

These are "typical" doses for these AAs.

AA

AA	DOSAGE
Cypro	5-12.5mg/day
Bica	50-100mg/day
Spiro	50-200mg/day

SEE DETAILED GUIDE FOR MORE INFO

4. QUICK METHOD GUIDE

After 3 months double estradiol doses. Maximal doses noted in estrogens. Starting on high levels of e2 is associated with slowed breast growth.

After a year, or reaching tanner 3-4, switch to injections (monotherapy dose) + progesterone (200mg rectal).

SEE DETAILED GUIDE FOR MORE INFO

5. BLOOD TESTS (LABS)

labs are useful as it's an objective measurement of your hormone levels. labs are also **very useful** in diagnosing issues and providing assistance.

Trough

Get labs at your trough. Trough means right before your next dose. Trough is important because you can gauge T suppression, your lowest E levels, and if you should adjust your dose.

Lab frequency

Get labs, when starting, at the 3mo mark. While adjusting your dose keep getting labs every 3mo. When your dose is refined, you can bump it to every 6mo-1yr.

Test for (always)

- Estradiol (e2)
- Testosterone (T)
- LH/FSH

Test for (conditional)

- Estrone (e1) - If: on oral e
- DHT - If: masculinization signs with suppressed T
- AST & ALT (Liver) - If: on bica or >=50mg cypro
- SHBG - If: goal: optimize free e2
- Prolactin - If: on cypro
- Potassium - If: on spiro

Liver toxicity is dangerous. It can lead to serious complications, and even death. That is, despite low incident rates for bica. If you're on a higher dose of cypro then liver levels tested as higher doses are associated with liver toxicity.

Target E levels

E2: E1 ratios: < 1.8 E2:E1 ratios
Free E: >= 1.5%

Target E2 monotherapy levels

200-300 pg/ml.
735-1,100 pmol/L
(1pg/mL = 3.671 pmol/L)

Target Total T levels

10-54 ng/dL.
0.341-1.87 nmol/L
(1ng/dL = 0.0347 nmol/L)

SEE DETAILED GUIDE FOR MORE INFO

6. ACQUIRING MEDS

The safest way is via your local pharmacy with a doctor's prescription, but this can be very difficult or impossible to option depending on your situation.

Lists of sources are provided in each thread. Since every country is different, e.g. varying strictness of customs, and sources constantly change, please read the thread and ask for help if you need it.

SEE DETAILED GUIDE FOR CUSTOMS INFO

7. STALLED BREAST GROWTH TIPS

Cycle oral e 2-6mg/daily ontop of usual regimen, 3mo on, and 3mo off. Tip is for when you're on e2 injections.

Cycle progesterone, 2 weeks on, and 2 weeks off (get on breasts and/or suppository).

You may alter these cycle lengths to your preference.

SEE DETAILED GUIDE FOR MORE INFO

8. FEMINIZING & PREVENTING BREAST GROWTH

Feminization without any breast growth cannot be done with 100% reliability.

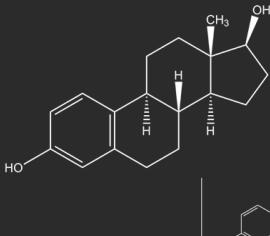
Raloxifene, a SERM, is commonly used to stunt, stall, or somewhat revert breast tissue growth. Raloxifene, in conjunction with e2, can be a competitive antagonist to e2 when it comes to breast growth (thus ralox dosage should out compete e2).

Methods

- Ralox 60-120mg/day (increases serum T/LH/FSH lvs).
- Ralox gel on breasts (2.5-5mg per breast, daily).
- DHT gel on breasts has direct anti-estrogenic action on breast cells and cannot be aromatized to e2 (like T).
- Cypro monotherapy has been done, but low sex hormones cause issues (bone mineral density and lacking neurosteroid action of sex hormones).
- Bica monotherapy has also been done, but heightened T levels will be aromatized into e2 (90% get gynecomastia).
- Do not weight cycle. Similarly, one would want to maintain weight/low healthy BMI. I.e. prevent/treat breast fat.

Regimens (examples)

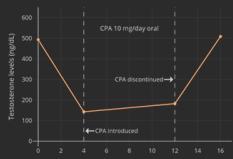
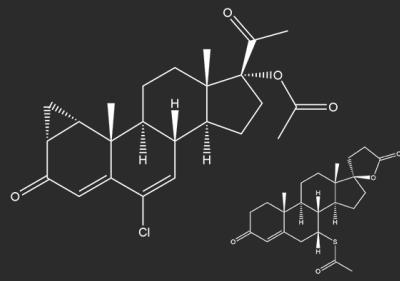
- 500mcg bca + 12.5mg cypro + 80mg ralox
- 100-150mcg bca + aromatase inhibitor + 80mg ralox
- GnRH + 50mg bca + 60mg ralox + low-mid dose e2



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1. ANTI-ANDROGEN (AA)

a) CYPROTERONE ACETATE (CYPRO, CPA)

Dosage: 10-20mg/day. 10-15mg/day will block female T levels to 40-50%. Studies have shown 10mg to be effective as an AA (50-70% lowered T @ 10mg/day). Action: Strong steroid AA, competes with aromatase. Nukes T production and weakly blocks T at the AR level (lowers T levels and blocks T). Due diligence: Doses above 12.5mg are not recommended as higher doses for prolonged periods are linked to venous thromboembolism, prostaticoma and meningioma (benign brain tumors). May raise prolactin levels. Monitor yourself for pseudo-lactation and monitor prolactin levels via blood tests. Higher doses of 100mg/day are associated with liver toxicity. In general prolactin is linked to vitamin B12 deficiency. To prevent B12 deficiency you should eat animal products, drink lots of milk, and/or take a B12 supplement to ensure healthy B12 levels.

Info: Effective at reducing erections and general function of gonads. Affordable AA.

b) BICALUTAMIDE (Bical)

Dosage: 25-100mg/day. 100-150mg/day will block female T levels which is achieved via estradiol at a sufficient dose. If not taking estradiol or if you have male levels you will need 50mg/day or more. 100-150mg/day at minimum appears to be needed to fully or nearly fully block 600ng/ml T.

Action: Silent non-steroidal AA antagonist that also blocks aromatase (i.e. weakly blocks T at the receptor level, and stops cells from being male). Does not lower T but it does blocks its effects with sufficient dosages. In conjunction with E bical increases T levels by 20-70%. Due to the blockage of the AR and T being utilized aromatization occurs (T > E) and may slightly raise E levels.

Due diligence: Bical has a low incidence rate of unfavorable liver level changes (3% at 150mg/day). Liver changes usually occur within the first 3 or 4 months of treatment, and therefore one should monitor liver levels in these first three months. It's advisable to get liver levels tested every 3-6mo. Liver toxicity (hepatotoxicity) is rare with bical, and is lower than the incidence rate of flutamide (.03% or 3 per 10,000). Avoid if you have a history of liver issues.

Info: Less effective at reducing general function of gonads. More expensive AA. Buildup time for bical to 50% of steady levels is reached after 1 week, about 80-90% steady state levels after 3-4 weeks, and 100% after 6-12 weeks of continuous daily administration.

c) SPIRONOLACTONE (Spiro)

Dosage: 50-200mg/day. At doses of 50-200mg/day spiro is more effective at blocking female T levels levels and less suited at blocking male T levels.

Action: A potassium sparing diuretic that is a weak AA, AR antagonist (blocks androgens) that also blocks aromatase synthesis (i.e. weakly blocks T from being produced). Variable efficacy at reducing T levels (ranges from nil to slight reduction). Main action is via being an AR antagonist (blocking the AR). It has a similar action as bical, but bical is far superior in this regard as per spiro.

Due diligence: Due to being a potassium sparing diuretic avoid potassium supplements. Risk factors for high potassium levels (hyperkalemia) includes old ages, kidney disease, and taking multiple potassium-elevating drugs, and intake of potassium supplements. In studies on children or young adults taking spiro potassium levels weren't significantly raised. At higher doses spiro use has been associated with elevated cortisol levels (associated with visceral adiposity), brain fog, and depression.

Info: Makes you pee a lot (somewhat mitigated via NSAIDs such as aspirin). Affordable AA. Start dose at <=100mg/day. Also used as an anti-cancer treatment and for hypertension (high blood pressure). Has off-target action with high antimarial/corticoid activity and mixed weak estrogenic and antiestrogenic or SERM-like activity (which has been implicated in having a negative impact on breast development, but effects are "weak" here so it's debatable).

d) GnRH Agonists (GnRH, GnRHa)

Types: Buserelin, leuprolide, goserelin, triptorelin.

Action: Agonist of the GnRH receptor which causes the pituitary gland to initially "spike" LH/FSH levels (LH causes luteinizing hormone to release T), and then to become desensitized to LH/FSH (lowering LH/FSH). It is like pressing the button the pituitary to constantly release LH/FSH. In response to these GnRH agonists, which causes it to "run out of fuel" and to become desensitized to GnRH. Thereby little to no LH/FSH is released with continuous usage because we're always pressing this button.

Info: Gold standard for suppressing LH/testosterone. Great alternatives if they can be obtained affordably, i.e. they are prohibitively expensive for most, and may be available through your GI/endo via insurance/public healthcare. Majority of "side effects" noted online are due to lacking sex hormones (E2/T/etc) and therefore don't apply w/HRT.

e) ESTRADIOL MONOTHERAPY (E MONO, E MONOTHERAPY)

Dosage: Estradiol (e2) levels of 200pg/ml suppress T levels by ~90%, and e2 levels between 200-500pg/ml suppress T levels by ~90-95%. This may vary due to capacity for gonadal to produce androgens (e.g. T), and therefore your "monotherapy levels" may differ here (e.g. 90% suppression of 400pg/ml is "sufficient" as compared to 90% suppression of 700pg/ml).

Action: Lowers LH/FSH via the brain regulating it has sex hormones, and via lowering LH/FSH it tells the gonads to stop producing T. In others, on the HPG axis it creates a negative feedback loop, lowers LH/FSH, and results in T production via the gonads significantly dropping with sufficient e2 levels (also lowers sperm production via lowering FSH levels).

Info: Easily and reliably attainable via injections. Can be attained via patches, gel, or sublingual.

2. ESTROGEN (e)

a) INJECTIONS

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

b) TRANSDERMAL

Types: Patches or gels. Patches come in reservoir or matrix forms. Reservoir patches are to be changed every 3-5 days. Matrix patches are to be changed every 7-8 days.

Dosing: Patching starts at 10-100pg/day depending on preferences and AA. 150-200pg/day or more is the final dose per your goal levels & labs. 100pg patch is approximately equal to 100pg/ml for levels. Transdermal gel starts at an equivalent dosage of 50-100pg patches. Gel with 0.06% concentration has 5mg estradiol in 2.5g gel. Estradiol levels achieved with 1.5mg of transdermal gel are similar to those with a 50ug patch, and 100ug patch is roughly equivalent to 3mg of gel. Gel must be applied daily.

Info: Patches are known to leave visible residue after removal. Reservoir patches are known to cause skin rashes (around 14-22% occurrence rate). Gel and transdermal application areas do change absorption, and smaller application areas for gel gives higher levels. Factors which may contribute to inter- and intraindividual variability with transdermal reservoirs include skin location and thickness, hair follicle density; solvent (alcohol) evaporation; skin dehydration, ambient temperature, and humidity; and reservoir size (for reservoir patches). Application area effectiveness: scrotum > buttocks > stomach > thigh = arm > hand = foot.

c) SUBLINGUAL (subS) & BUCCAL

Types: Estradiol hemihydrate (17-beta estradiol) or estradiol valerate (EV) pills can be used.

Dosing: Start at 2mg/day and later increase to 4-8mg/day. Split doses throughout the day as to maintain steadier levels. 2mg dosing example: 0.5mg/4x day (e.g. 16hr day: every 4 hrs take 0.5mg). It is recommended to split your dose >=3/4 for more stable levels (single-dose oral e2 half-life ranges from 2-20hrs for 0.5mg-1mg).

Info: Sublingual administration method is to dissolve under your tongue, and buccal is to dissolve between your cheek and gums. Do not eat or drink while they are dissolving, and it is a general rule of thumb to wait 10-15min before eating or drinking after the pills have dissolved. Splitting or crushing the pills may help them dissolve faster, and this is especially applicable to pills such as progynova (EV) due to its sugar coating. Sub e has an estro (e1) spike ~2hrs after administration, and it has been speculated that this is the result in estradiol being taken up by the reticuloendothelial system and then metabolized into estro (e1). It is not via "accidental swallowing" primarily.

d) ORAL

Types: Estradiol hemihydrate (17-beta estradiol) or estradiol valerate (EV) pills can be used. 2mg EV is roughly equivalent to 1.5mg estradiol hemihydrate.

Dosing: Start at 2mg/day and later increase to 6-8mg/day. Split doses throughout the day as to maintain steadier levels. 2mg dosing example: 0.5mg/4x day (e.g. 16hr day: every 4 hrs take 0.5mg). It is recommended to split your dose >=3/4 for more stable levels (single-dose oral e2 half-life ranges from 12-20hrs for 0.5mg-1mg).

Info: Oral administration method is to swallow the tablets. Significantly raises SHBG levels (sex hormones, such as e2, bind to SHBG which makes them "inactive" and not "free hormones"), and thus 5-10mg/day bical is recommended as to counteract this. Due to lower e2 levels the main consideration when starting on oral e is T suppression. Cypro is more reliable here, and for bical it's recommended to get a pre-hrt blood test (for T) as to determine the proper dosage as to block.

e) CREAMS

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

f) TOPICAL

Types: Estradiol valerate (EV) is cheap and widely available via prescription.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

g) INHALATION

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

h) INTRAVAGINAL

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

i) INTRAMUSCULAR

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

Dosing: Adjust dose as needed based on blood tests. EV is injected at once a week or less. EC & EE are better if you want to go between two weeks between injections. Make sure to check the concentration on the vials. If the concentration is 40mg/ml, then you dose 0.25ml, to get 10mg of estradiol. Utilize the "injectable estradiol simulator" (search it) for determining dose and injection frequency (cycle length). With a 250pg/ml trough for estradiol monotherapy dosages are: 5mg/5days or 9mg/week for EV, 5mg/week for EC and 4mg/week for EE.

Info: Can be done intramuscularly (IM) or subcutaneously (subQ, SQ). Similar efficacy, potency, and levels for each administration method for EC (per studies).

j) SUBCUTANEOUS

Types: Estradiol valerate (EV) is cheap and widely available via prescription. It has a higher and faster initial peak that quickly drops off. Estradiol cypionate (EC) is more expensive, has a lower initial peak, but it lasts longer and gives steadier levels. Estradiol enanthate (EE) is the longest lasting and is considered the best for injecting every two weeks as it raises slightly higher than EC, has slightly higher overall levels, and is similarly steady for longer.

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k) INTRADERMAL

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m) INTRAMAMMARY

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o) INTRAMUSCULAR

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